



DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Community Living

Agency Information Collection Activities; Proposed Collection; Comment Request;

Alzheimer's and Dementia Program Data Reporting Tool (ADP-DRT) OMB Control

Number 0985-0022

AGENCY: Administration for Community Living, HHS.

ACTION: Notice.

SUMMARY: The Administration for Community Living (ACL) is announcing an opportunity for the public to comment on the proposed collection of information listed above. Under the Paperwork Reduction Act of 1995 (the PRA), Federal agencies are required to publish a notice in the *Federal Register* concerning each proposed collection of information, including each proposed extension of an existing collection of information, and to allow 60 days for public comment in response to the notice. This notice solicits comments on the proposed Revision for the information collection requirements related to Alzheimer's and Dementia Program Data Reporting Tool (ADP-DRT).

DATES: Comments on the collection of information must be submitted electronically by 11:59 pm (EST) or postmarked by [PLEASE INSERT 60 DAYS FROM THE DATE OF PUBLICATION IN THE *FEDERAL REGISTER*].

ADDRESSES: Submit electronic comments on the collection of information to: Erin Long (erin.long@acl.hhs.gov). Address written comments on the collection of information to Administration for Community Living, Washington, D.C. 20201, Attention: Erin Long PRA comments Alzheimer's and Dementia Program Data Reporting Tool (ADP-DRT)

FOR FURTHER INFORMATION CONTACT: Erin Long, erin.long@acl.hhs.gov, 202-795-7389.

SUPPLEMENTARY INFORMATION: Under the PRA (44 U.S.C. 3501-3520), Federal agencies must obtain approval from the Office of Management and Budget (OMB) for each collection of information they conduct or sponsor. “Collection of information” is defined as and includes agency requests or requirements that members of the public submit reports, keep records, or provide information to a third party. The PRA requires Federal agencies to provide a 60-day notice in the *Federal Register* concerning each proposed collection of information, including each proposed extension of an existing collection of information, before submitting the collection to OMB for approval. To comply with this requirement, ACL is publishing a notice of the proposed collection of information set forth in this document. With respect to the following collection of information, ACL invites comments on our burden estimates or any other aspect of this collection of information, including:

- (1) whether the proposed collection of information is necessary for the proper performance of ACL's functions, including whether the information will have practical utility;
 - (2) ways to enhance the quality, utility, and clarity of the information to be collected;
 - (3) accuracy of ACL's estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used to determine burden estimates;
- And (4) ways to minimize the burden of the collection of information on respondents, including through the use of automated collection techniques when appropriate, and other forms of information technology.

The Older American’s Act requires ACL to evaluate “demonstration projects that support the objectives of this Act, including activities to bring effective demonstration projects to scale with a prioritization of projects that address the needs of underserved populations, and promote partnerships among aging services, community-based organizations, and Medicare and Medicaid providers, plans, and health (including public health) systems. (Section 201 (42 U.S.C. 3011) Sec. 127. Research and Evaluation).

To fulfill the evaluation requirements and allow for optimal federal and state-level management of ACL’s Alzheimer’s Disease Program, specific information must be collected from grantees. The current reporting tool is set to expire 12/31/2023. The Alzheimer’s and Dementia Program (ADP) Project Officer has reviewed the current data collection procedures to ensure the acceptability of these items as appropriate and thorough evaluation of the program, while minimizing burden for grantees.

The result of this process is the proposed modifications to the existing data collection tool. ACL is aware that different grantees have different data collection capabilities. It is understood that, following the approval of the modified data collection tool, ACL will work with its grantees to offer regular training to ensure minimal burden.

To support alignment with Executive Order 13985 on Advancing Racial Equity and Support for Underserved Communities Through the Federal Government, Executive Order 14075 on Advancing Equality for Lesbian, Gay, Bisexual, Transgender, Queer, and Intersex Individuals, and Executive Order 13988 on Preventing and Combating Discrimination on the Basis of Gender Identity and Sexual Orientation, ACL is adding three sexual orientation and gender identity (SOGI) items to the ADP-DRT. Understanding these disparities can and should lead to improved service delivery for ACL’s programs and populations served.

The proposed data collection tools may be found on the ACL website for review at <https://www.acl.gov/about-acl/public-input>.

ESTIMATED PROGRAM BURDEN:

ACL estimates the burden associated with this collection of information as follows:

Type of Respondent	Form Name	Number of Respondents	Frequency of Response	Average Time per Response (in hours)	Total Burden Hours (Annual)
Grantee	ADSSP-DRT	69	2	6.64	916.32
Total					916.32

Dated: October 18, 2023.

Alison Barkoff,

Senior official performing the duties of the Administrator and the Assistant Secretary for Aging.

[FR Doc. 2023-23417 Filed: 10/23/2023 8:45 am; Publication Date: 10/24/2023]